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(54) Title: PROCESS FOR THE PREPARATION OF A MELAMINE BASED FLAME RETARDANT AND POLYMER COM-**POSITION**

(57) Abstract: The invention relates to a process for the preparation of a melamine based flame retardant by a reaction of a melamine comprising compound and a polyol in a molar ratio between 1.0: 1 and 4.0: 1, whereby the reaction is performed in an extruder equipped with a degassing means, at a temperature between 200 and 300°C. Furthermore the invention relates to a melamine based flame retardant obtainable by said process and a polymer composition comprising said melamine based flame retardant.



WO 2004/055029 PCT/EP2003/050955

-1-

Process for the preparation of a melamine based flame retardant and polymer composition

The invention relates to a process for the preparation of a melamine based flame retardant by a reaction in an extruder between a melamine comprising compound and a polyol with the optional addition of a polymeric carrier material.

Such process is known from *US-A-4,010,137*, which discloses a process wherein melamine pyrophosphate and pentaerythritol are combined as single component flame retardants and heated in a vessel at 175°C – 275°C for 0.5 to 4 hours. Specific degassing conditions insure the incorporation of an effective amount of the flame retardant into the synthetic resin. No significant foaming is observed when the synthetic resin composition is subjected to moulding conditions. The resulting product is used especially for the preparation of flame retardants containing polypropylene. Furthermore, this publication teaches the need for stable melamine based flame retardants which can be moulded at temperatures up to 260°C. The reaction product of H₃PO₄, P₂O₅, dipentaerythritol and melamine is disclosed as a particularly stable flame retardant. This product can be moulded without foaming at a temperature up to 246°C. A clear disadvantage of the process as disclosed in this reference results from the fact that the preparation of the flame retardant requires a process time of at least 0.5 hours and up to 4 hours.

A process for the preparation of a melamine based flame retardant is disclosed in WO 00/68337. This reference discloses a process wherein powder blends comprising pentaerythritol and melamine phosphate are prepared. These powder blends, however, are strongly limited in their applicability for the manufacture of polymer compositions, which is caused by the fact that foaming occurs during the processing of the powder blend in the polymer, e.g. during the compounding or injection moulding.

A severe problem results from the fact that single component flame retardants that contain phosphoric ester groups are subject to hydrolysis. In the event that these flame retardant additives are blended with polymer materials, such as polypropylene, their moisture sensitivity causes considerable leaching of the flame retardant molecules from the polymer matrix resulting in reduced flame retardant efficiency of the polymer composition and decreased electrical insulation properties.

Various methods are known to increase the water resistance of polymeric flame retardant compositions. They include partial phosphorylation of polymers, such as polyvinyl acetate, encapsulation of polar flame retardant components, such as acid source APP particles with copolymer of vinyl pyrrolidone and comonomer, modification of flame retardants with surfactants (non-ionic or ionic) or the replacement of polyols (polar species) by other char

formers, such as dialkyl tin oxide, dialkyl tin dialkoxide, or polyol(alkylcarbonate). Some of these methods are effective in improving the water resistance of the polymeric flame retardant compositions, but result in complex additional processes often accompanied by environmental processes.

An object of the present invention is to provide a faster process for the preparation of a melamine based flame retardant and its master batch with high flowability.

Another object of the invention is the preparation of polymeric flame retardant compositions of increased water resistance while preserving the flame retardant performance (UL94 V-0) and mechanical properties.

These objects are achieved by the process according to the present invention, which relates to a process for the preparation of a melamine based flame retardant by a reaction of a melamine comprising compound and a polyol.

The melamine based flame retardant obtainable by the process of the present invention is particularly suitable for producing flame retardant polymer compositions of high water resistance. Due to the higher thermal stability of the melamine-based flame retardant obtainable by the process of the present invention, the polymer composition can be moulded at higher temperatures as compared to the polymer compositions disclosed in *US-A-4,010,137*.

A flame retardant polymer composition comprising a mixture of a melamine comprising compound and pentaerythritol is disclosed in *US-A-4*,632,946. This reference only discloses polymer compositions that can be processed at melt temperatures up to 230°C, whereas at higher temperatures foaming will occur. This confirms the stability problem as already mentioned in *US-A-4*,010,137.

A further advantage of the process of the invention is established by the fact that the melamine based flame retardant obtained exhibits a higher degree of stability than the melamine based flame retardant obtained according to the process as described in *US-A-4,010,137*. A composition comprising the flame retardant of Embodiment II of *US-A-4,010,137* has to be moulded below the temperature 232°C, in order to prevent foaming. The melamine-based flame retardant according to the process of the present invention does not show any foaming at temperatures up to 280°C.

The invention, therefore, also relates to a melamine based flame retardant obtainable by the process of the present invention.

The process according to the present invention that relates to a process for the preparation of a melamine based flame retardant by a reaction of a melamine comprising compound and

a polyol is characterised in that the melamine comprising compound is selected from the group consisting of melamine phosphate, melamine pyrophosphate and melamine polyphosphate, the polyol is selected from the group consisting of pentaerythritol, dipentaerythritol and tripentaerythritol and the reaction is carried out by reactive extrusion in an extruder in a molar ratio of the melamine comprising compound to the polyol between 1.0:1.0 and 4.0:1.0 and that the reaction is performed at a temperature between 200 and 300°C.

In a preferred embodiment of the invention the melamine based flame retardant is prepared by reaction of melamine phosphate with pentaerythritol.

With the process of the invention a faster process for the preparation of the melamine based flame retardant is obtained within a short time period.

The melamine comprising compound, such as melamine phosphate used in the process according to the invention and the polyol, such as pentaerythritol, can be of any standard chemical quality. Preferably they are used in the form of a powder. Pentaerythritol in pure form melts at 260°C and in mixtures below 260°C. Preferably fine powder forms are used. The melamine phosphate or the other melamine comprising compounds and the pentaerythritol as well as the other polyols can be fed separately to the extruder or can be premixed with e.g. a high shear mixer. Preferably a premixed mixture is used. This reduces the necessity of the control of the amount of flows of the material.

In the process of the invention the melamine comprising compound, such as melamine phosphate, and the polyol, such as pentaerythritol, are reacted in an extruder. This process embodiment is referred to as reactive extrusion. Extruders, such as the ones routinely used in the polymer processing industry, e.g. mono or double screw extruders, are suitable for this purpose. A twin-screw extruder is particularly preferred for the handling of fine powders. In the process of the invention the extruder is equipped with a degassing device. Generally this degassing means is a port or opening in the extruder that is connected to a vacuum pump.

In the extruder the melamine-based flame retardant is produced, which leaves the extruder through the die as an extrudate. The extrudate is cooled and shaped, preferably by cutting, into formed particles, preferably pellets.

In the process of the invention the melamine comprising compound and the polyol react to form a melamine phosphate-polyol condensate, whereby corresponding molar equivalents of water are released. The water leaves the extruder through the degassing device. In this application the melamine phosphate-pentaerythritol condensate is also referred to as the melamine based flame retardant.

The molar ratio of the melamine phosphate to pentaerythritol is in the range between about 1.0: 1.0 and 4.0: 1.0. Preferably the molar ratio is chosen between about 2.0: 1.0 and 2.4: 1.0. Melamine based flame retardants of excellent stability are obtained. Most preferably, the molar ratio is about 2.2: 1.0. In the context of the description of the invention thermal stability is defined as the degree of resistance against foaming upon heating of the melamine based flame retardant. For a more precise differentiation in thermal stability of flame retardant compositions physicochemical methods, such as thermo gravimetric analysis (TGA) and differential scanning calorimetry (DSC), can be used.

The reaction of the melamine comprising compound and the polyol to obtain the melamine based flame retardant can be performed at temperatures between about 100°C and 300°C. However, for a complete conversion the reaction must be performed at a temperature higher than 200°C.

The maximum temperature for the reaction is chosen below 300°C. Preferably, the reaction of melamine phosphate and pentaerythritol is carried out in a temperature range between 220°C and 280°C. Between 220°C and 280°C a good balance is obtained between the rate of reaction and the degradation of the reaction product melamine based flame retardant. More preferably, the reaction of the melamine phosphate and the pentaerythritol is conducted at a temperature between 240°C and 260°C. The thermal stability of the melamine based flame retardant produced in this temperature range is excellent.

The time period for reaction in the process of the present invention is, in general, between 1 and 20 minutes. It was found that 1 batch in the extruder at the particularly preferable temperature range of 240 – 260°C is sufficient for producing a non-foaming melamine based flame retardant. In an alternative embodiment of the process, the extruder is fed once again with the melamine based flame retardant obtained. With 2 passages through the extruder the doubling of the residence and hence reaction time is achieved. Due to a longer reaction time period in the extruder an increased stability of the melamine based flame retardant obtained can be achieved. This difference in stability can be easily assessed by TGA, which method is described below. Furthermore, the time period for reaction can also be changed with other means, e.g. by adjusting the number of revolutions per minute of a screw in the extruder, or by using a so-called static mixer in the extruder. Foaming of the reaction product can be determined by applying the foaming test as described below.

It has been found advantageous to add polymer material to the extruder as a carrier resin in addition to the melamine comprising compound and the polyol. A more constant extruder output is achieved with less than 30 weight% of polymer present in the flame retardant composition, especially when the extruder operates at temperatures below 270°C. The

- 5 -

amount of the polymer in general is kept low, e.g. between 5.0 - 30.0 weight, particularly between 5.0 - 20.0%, related to the total weight amount of the flame retardant composition. When exposed to water a flame retardant composition in the presence of suitable polymer material, such as polypropylene, has a higher resistance to water, which is desirable in protecting polymer material against erosion caused by the so-called leaching effect of the flame protecting agent, which in this case is the reaction product of melamine comprising compound phosphate and the polyol.

In a preferred embodiment of the invention the reaction of the melamine comprising compound and the polyol is carried out in the presence of a third component, particularly 5.0 to 20.0 weight% of a polymer.

In a particularly preferred embodiment of the invention the reaction of the melamine phosphate and pentaerythritol is carried out in the presence of 5.0 to 20.0 weight% of a polymer.

Any type of polymer material can be chosen that is suitable for melt processing at the extruder temperature, preferably at processing temperatures below 300°C. In general the polymer or carrier resin is chosen according to the polymer matrix material that needs flame retardation. Polyethylene is the first choice due to its large availability and easy processing properties. It has been found in this respect that through the use of high density polyethylene, HDPE, light coloured melamine based flame retardant pellets can be produced, which is advantageous for producing light coloured, flame retardant polymer compositions. In an alternative embodiment the use of polypropylene has been found advantageous. An acceptable colour of the flame retardant master batches is obtained, combined with high fluidity and excellent flame retardancy and mechanical properties of the composite material. Furthermore through the use of a polymer it is easier to obtain pellets of the melamine based flame retardant.

Other polymers suitable for the polymer composition of the present invention are those polymers, which are processed at temperatures below 300°C and preferably below 280°C. An example of these polymers is a polyolefin, e.g. polypropylene (PP), either as homo-, co-or terpolymer.

A flame retardant composition comprising polypropylene as polymer material has a good thermal stability during processing. Its decomposing temperature is in agreement with the pyrolysis point of PP. No typical melt process, but a softening zone above 180°C is observed. The shaped particles can deform and flow when compounding with PP and easily obtain an effective blending and dispersing. In compounding the softening flame retardant composition can plasticise the PP resin melt. A small amount of PP can also improve the

flowability of the composition considerably. Therefore, a master batch product of flame retardant content as well as high flowability is obtained. The flame retardant particles can deform and arrange in the flow direction of the polymer melt to form a typical fiber structure.

A non-exhaustive list of additional polymer materials is listed here below:

Polymers containing styrene, such as HIPS (High Impact polystyrene), polyamides, polyimides, polyesters, polycarbonates, epoxy resins, polyurethanes and mixtures and blends of these materials.

The polymer composition according to the invention can be produced by known techniques as e.g. extrusion, compounding or injection moulding. The amount of the melamine based flame retardant obtainable by the process of the present invention in the polymer composition depends on the required level of flame retardancy. This required level of flame retardancy varies per application. Generally, for electronic or construction applications different amounts are required. An amount of up to 30 weight% is sufficient for assuring flame retardancy in many applications. The flame retardant composition may comprise other additives or auxiliary materials to improve the properties of the composition against decomposition, discolouration or oxidation, e.g. colorants, pigments, thermal and UV-stabilisers, mould release agents, co-flame retardants, softeners, lubricants, antioxidants, flow-promoting agents, dispersing agents, fillers and reinforcing agents as normally used in the formulation of polymer compositions.

Examples of coating compositions are pigment-filled alkyd resins, homo- or copolymers based on acrylates, urethanes, esters, epoxies or modifications thereof. Optionally these can be provided with the usual cross linkers. After curing of this coating composition a coating is formed. In those processes where the flame retardant compositions are incorporated into a liquid polymer, a polymer solution or a polymer dispersion, for example a coating composition, the choice of auxiliary material is determined by the solubility of the auxiliary material in said liquid polymer, polymer solution or polymer dispersion.

Suitable organic auxiliary materials are organic compounds, polymers or copolymers based on vinylpyrrolidone, vinyl acetate and vinyl caprolactam, or mixtures thereof. Also suitable are the above-mentioned polymers or copolymers or additional polymers or copolymers based on epoxides, urethanes, acrylates, esters, amides, stearates, olefins, cellulose derivatives or mixtures thereof. In the event that the compositions are prepared from aqueous slurry, water-soluble organic auxiliary materials are of advantage because they can easily be added to this slurry.

When the melamine based flame retardant obtainable by the process of the present invention is added to a liquid polymer, a polymer solution or a polymer dispersion, which contains water, the agglomerate is easily dispersible. Polyvinyl pyrrolidone, polyvinyl alcohol and polyvinyl caprolactam are easy to handle and can be used in a wide range of applications due to their good solubility in water.

The quantity of organic auxiliary material amounts to 0.1 to 10.0 wt%, relative to the total weight of the composition. If very high flame retardation requirements are to be met with the melamine based flame retardant obtainable by the process of the present invention, preferably 0.1 to 5.0 wt% is used, even more preferably 0.1-3.0 wt% relative to the total agglomerate.

It has been found that the advantages mentioned for melamine based flame retardant obtainable by the process of the present invention also apply to compositions of other flame retardant compounds, including halogen containing and halogen-free ones. However, preference is given to halogen-free flame retardant compounds, including triazine compounds, such as melamine, ammeline and/or ammelide, higher condensation products thereof, such as melam, melem and or melon, melamine derivatives, such as melamine salts, e.g. melamine phosphate salts, melamine acetate, melamine pyrophosphate salts, melamine polyphosphate and/or melamine ammonium polyphosphate salts; metal compounds, such as aluminium hydroxide, magnesium hydroxide, antimony trioxide, Sb₂O₅, zinc oxide, sodium antimonate, zinc stannate and/or zinc borate.

The flame retardant properties of the flame retardant composition as obtained by the process according to the present invention can be enhanced by the presence of additional compounds exerting a synergistic effect with the flame retardant composition, such as a charforming compound, optionally combined with a catalyst promoting the formation of char in the polymer composition. In general, any compounds capable of enhancing the flame-retardant properties by means of char formation are suitable, such as resinous polymers selected from the group consisting of phenolic, epoxy, polyurethane, novolak, melamine, alkyd, allyl, unsaturated polyester, silicone, urethane and acrylate resins, polyphenylene ether, polycarbonate, polyvinyl alcohol, poly(ethylene covinyl), starch or glucose.

Metal salts of tungstic acid, a complex oxide acid of tungsten and metalloids, salts of tin oxide, ammonium sulphamate and/or its dimer may be used as catalysts for promoting char formation.

The invention are further illustrated with the following, non-limiting examples and Comparative Experiments.

PCT/EP2003/050955 WO 2004/055029

-8-

In the application examples reference is made to various test methods, which are explained below:

- LOI (Limiting Oxygen Index) value: is measured using an ATLAS Limiting Oxygen Index instrument on sheets of 120 mm (I) x 6.5 mm (b) x 3 mm (t) according to ASTM D2863-77. A higher LOI value indicates better flame retardancy;
- Flammability test is measured according the norm UL 94 HB. UL 94 contains the test procedure for the horizontally positioned solid plastic specimen in the form of rods. The dimensions of the rods are 127 mm (I) x 12.7 mm (b). The thickness(t) of the rods should correspond to the minimum for the Intended application. Tensile test and elongation at break is performed according to ISO 527-1A;
- Impact test: the notched impact strength is measured using a XJ-40A Impact strength tester according to ISO 180 -A/U;
- Foaming test: is used for fast simulation of moulding and check of the stability of a product. In the foaming test an amount of sample is put in a beaker that is placed in an oven at a temperature of 280 °C for 30 minutes. Foaming is visually evaluated;
- TGA: thermo gravimetric analysis is performed with a Dupont 2100 thermal analyser. Samples are heated from room temperature at a rate of 20 °C/min, whereby the weight loss versus temperature is recorded.

Example !: Preparation of a melamine based flame retardant using an extruder.

Melamine phosphate and pentaerythritol are premixed in a high shear mixer at a molar ratio of 2.2: 1.0. The premix obtained is fed to a twin-screw extruder (o=30 mm; L/D=32; fitted with a vacuum degassing), which operates at a screw speed of 35 rpm. Additionally HDPE is fed to the extruder in a HDPE: premix ratio of 5: 20. The average temperature in the extruder is 250°C and the residence time is between 3 and 5 minutes, on average 4 minutes. The extrudate obtained is cut into pellets and subsequently dried. The pellets have a light colour. The stability of the pellets is determined with the foaming test and TGA. The results are summarised in Table 1. It can be seen that the melamine-based flame retardant is very stable since it does not show foaming at 280°C.

Examples II-IV: Preparation of melamine based flame retardants, by several passes through an extruder.

The melamine based flame retardant as obtained in Example I is again fed to the twin-screw extruder and given the same treatment as described in Example I. A longer reaction time is

WO 2004/055029

-9-

obtained, which amounts to up to 16 minutes by passing the extruder 4 times. The following residence times are obtained:

Example II: 2 extruder passes: total residence time = 8 minutes

Example III: 3 extruder passes: total residence time = 12 minutes

Example IV: 4 extruder passes: total residence time = 16 minutes

The stability of the pellets is determined by applying the foaming test and TGA. The results are summarised in Table 1. As in Example I the flame retardants as obtained according to these examples do not show foaming. Furthermore the TGA results show a lower weight loss of the product of Example IV, which reflects that the stability of the flame retardant is further increased upon longer total residence time.

Comparative Experiments A. B: Heating of melamine pyrophosphate and pentaerythritol at 250°C, according to Embodiment II of Table I in US-A-4,010,137.

Melamine pyrophosphate and pentaerythritol are premixed in a high shear mixer at a molar ratio of 2.2; 1.0. The premix obtained is placed in a beaker that is heated in an oven at 250°C. The heating is conducted for 20 minutes (Comparative Experiment A) or for 40 minutes (Comparative Experiment B).

The stability of the obtained product is determined by applying the foaming test. The results are summarised in Table 1.

Table 1: Stability Tests of melamine based flame retardants

Melamine based flame retardant	Reaction time at 250°C	Foaming?	TGA* [wt%]	
Obtained according to	[minutes]			
Example I	4	No	7	
Example II	8	No		
Example III	12	No		
Example IV	16	No	5	
Comparative Experiment A	20	Yes		
Comparative Experiment B	40	Yes		

^{*}Weight loss in weight% at 300°C

Example V: The experimental conditions and procedures are the same as the ones applied in Example I except that the molar ratio of melamine phosphate and pentaerythritol is 1.6: 1.0 and the additional carrier resin is polypropylene (PP) with a PP: premix ratio of 10: 90.

Example VI: The experimental conditions and procedures are the same as the ones applied in Example V except that the additional carrier resin is high HIPS with a HIPS: premix ratio of 10:90.

Examples VII-IX: Preparation of a polymer composition comprising the melamine-based flame retardant according to the invention.

A polymer composition based on polypropylene (PP, homopolymer, melt index 3.4 g/10 min [according to ISO 1133 (Procedure A) at 230°C, 2.16 kg] comprising the melamine based flame retardant as obtained according to Example V-VI and two anti-oxidants, is produced. The composition as defined in Table 2 is fed to a twin-screw extruder as described in Example I, which operates at 70 rpm and at a temperature of 190°C. The extrudate leaving the extruder is cooled in a water bath, cut into pellets and dried. The following polymer compositions are obtained:

Example VII: The flame retardant composition is prepared according to Example I.

Example VIII: The flame retardant composition is prepared according to Example V.

Example IX: The flame retardant polymer matrix is HIPS and the flame retardant used is prepared according to Example VI.

From these pellets test specimen are moulded at a temperature of 250°C. The moulding is performed without problems. No foaming or entrapped gas is found in the test specimen. Some properties of the specimen are evaluated and are summarised in Table 2. The tensile strength and impact strength is at the level of pure PP that has tensile strength and impact strength of 36.1 MPa and 45 l/m, respectively. Furthermore the LOI is increased from 17.5 for pure PP to 28 and the UL 94 level is increased from failure to V-O rating for the polymer composition.

<u>Comparative Experiment C:</u> Preparation of a polymer composition comprising a mixture of melamine phosphate and pentaerythritol.

A polymer composition is produced in a manner analogous to Example VII. As flame retardant a powder mixture of melamine phosphate and pentaerythritol (ratio 2.2: 1.0) is fed to the extruder together with the polypropylene. The test specimen is moulded analogous to the procedure as summarised in Example VII. During moulding foaming occurred. Furthermore, - 11 -

the mechanical properties are reduced, as can be seen in Table 2, as compared to pure PP that has a tensile strength of 36 MPa.

Table 2: Polymer composition comprising melamine based flame retardant

Composition	Example	Example	Example	Comparative
	VII	Vill	IX	Experiment C
PP Polymer	74.4	77.2		79.4
HIPS polymer			66.1	
Reaction product melamine	20.0	20.0	30.0	
phosphate and penta-			Į	
erythritol				
Carrier Resin	5.0 (Poly-	2.2 (Poly-	3.3 (HIPS)	
	ethylene)	propylene)		
Mixture melamine phos-				20
phate and pentaerythritol		 	į.	j
Dilauryl thiodipropionate	0.3	0.3	0.3	0.3
Bis(2.4-dicumylphenyl pen-	0.3	0.3	0.3	0.3
taerythritoldiphosphate				
Tensile Strength ¹⁾ [Mpa]	35.4	31.0		27.7
Impact Strength ²⁾ [J/m]		44.0		
Elongation at Break ³⁾ [%]	6.1	100.0		2.8
LOI ⁴⁾ [%]	23	28	27	24
UL 94 3.2 mm	Failure	V-0		Failure

For pure PP polymer: 1)36.1 Mpa, 2)45 J/m, 3)410%, 4)17.5%, 5)Failure

Claims:

- 1. A process for the preparation of a melamine based flame retardant by a reaction of a melamine comprising compound and a polyol, characterised in that the melamine comprising compound is selected from the group consisting of melamine phosphate, melamine pyrophosphate and melamine polyphosphate, the polyol is selected from the group consisting of pentaerythritol, dipentaerythritol and tripentaerythritol and the reaction is carried out by reactive extrusion in an extruder in a molar ratio of the melamine comprising compound to the polyol is between 1.0 : 1.0 and 4.0 : 1.0 and that the reaction is performed at a temperature between 200 and 300°C.
- 2. A process according to claim 1, characterised in that the melamine based flame retardant is prepared by reaction of melamine phosphate with pentaerythritol.
- 3. A process according to claim 1, characterised in that the extruder is a twin-screw extruder.
- 4. A process according to any one of claims 1–3, characterised in that the reaction of the melamine comprising compound and the polyol is carried out in the presence of up to 30 weight% of a polymer.
- A process according to claim 1, characterised in that the reaction of the melamine comprising compound and the polyol is carried out in the presence of 5 to 20 weight% of a polymer.
- 6. A process according to claim 4, characterised in that the polymer is selected from the group consisting of polyethylene, polypropylene and high impact polystyrene.
- 7. A process according to claim 6, characterised in that the polyethylene is high-density polyethylene.
- 8. The melamine based flame retardant obtainable by the process according to claim 1.
- 9. A polymer composition comprising at least a polymer and up to 30 weight% of the melamine based flame retardant according to claim 8.
- A polymer composition according to claim 9, characterised in that the polymer is polypropylene.
- 11. Shaped articles comprising the polymer composition according to claim 8.

INTERNATIONAL SEARCH REPORT

PCT/EP 03/50955

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 C07F9/02 C09K21/12 C08K5/3492 According to International Patent Classification (IPC) or to both national classification and IPC B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) C09K C07F C08K Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the International search (name of data base and, where practical, search terms used) EPO-Internal, WPI Data, PAJ C. DCCUMENTS CONSIDERED TO BE RELEVANT Relevant to claim No. Citation of document, with indication, where appropriate, of the relevant passages 8-11 X HALPERN Y ET AL: "Fire retardancy of thermoplastic materials by intumescence" INDUSTRIAL & ENGINEERING CHEMISTRY RESEARCH, AMERICAN CHEMICAL SOCIETY. WASHINGTON, US, vol. 23, no. 2, 1984, pages 233-238, XP000960807 ISSN: 0888-5885 Scheme II, Formula (3) page 235, at least 20% of (3) used in polypropylene US 4 010 137 A (BRADY DONNIE G) 1 - 11A 1 March 1977 (1977-03-01) column 4, lines 25-27 claim 1 table 1 -/--Patent family members are listed in annex. Х Further documents are listed in the continuation of box C. Special categories of cited documents: 'T' later document published after the international filing date or priority date and not in conflict with the application but died to understand the principle or theory underlying the "A" document defining the general state of the art which is not considered to be of particular relevance invention "E" earlier document but published on or after the international "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such docu-"O" document referring to an oral disclosure, use, exhibition or ments, such combination being obvious to a person skilled in the art. document published prior to the international filing date but later than the priority date claimed "&" document member of the same patent family Date of the actual completion of the international search Date of mailing of the international search report 10/08/2004 3 August 2004 Name and mailing address of the ISA Authorized officer European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Marquis, D Fax: (+31-70) 340-3016

INTERNATIONAL SEARCH REPORT

Internation Application No PCT/EP 03/50955

		PCT/EP 03/50955			
C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT					
ategory °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.			
A	US 4 140 856 A (BOST HOWARD W ET AL) 20 February 1979 (1979-02-20) example 1	1-11			
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INTERNATIONAL SEARCH REPORT

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Internation Application No
PCT/EP 03/50955

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
US 4010137	Α	01-03-1977	US	3936416 A	03-02-1976
US 4140856	Α	20-02-1979	NONE		

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